PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter I of the Patent Cooperation Treaty)

(PCT Rule 44bis)

Applicant's or agent's file reference 152528-280	FOR FURTHER ACTION	See item 4 below		
International application No. PCT/JP2005/003787	International filing date (day/month/year) 04 March 2005 (04.03.2005)	Priority date (day/month/year) 04 March 2004 (04.03.2004)		
International Patent Classification (8th edition unless older edition indicated) See relevant information in Form PCT/ISA/237				
Applicant MEIJI SEIKA KAISHA, LTD.				

1.	This international preliminary r International Searching Author	report on patentability (Chapter I) is issued by the International Bureau on behalf of the ity under Rule 44 bis.1(a).	
2.	This REPORT consists of a total of 7 sheets, including this cover sheet. In the attached sheets, any reference to the written opinion of the International Searching Authority should be read as a reference		
··	to the international preliminary	report on patentability (Chapter I) instead.	
3.	This report contains indications	relating to the following items:	
	Box No. I	Basis of the report	
	Box No. II	Priority	
	Box No. III	Non-establishment of opinion with regard to novelty, inventive step and industrial applicability	
	Box No. IV	Lack of unity of invention	
	Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement	
	Box No. VI	Certain documents cited	
	Box No. VII	Certain defects in the international application	
	Box No. VIII	Certain observations on the international application	
4.		ommunicate this report to designated Offices in accordance with Rules 44bis.3(c) and 93bis.1 but makes an express request under Article 23(2), before the expiration of 30 months from the priority	

	Date of issuance of this report 29 November 2006 (29.11.2006)
The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland	Authorized officer Masashi Honda
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Form PCT/IB/373 (January 2004)

PATENT COOPERATION TREATY

TRANSLATION From the INTERNATIONAL SEARCHING AUTHORITY To: **PCT** WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY (PCT Rule 43bis.1) Date of mailing (day/month/year) Applicant's or agent's file reference FOR FURTHER ACTION 152528-280 See paragraph 2 below International application No. International filing date (day/month/year) Priority date (day/month/year) PCT/JP2005/003787 04.03.2005 04.03.2004 International Patent Classification (IPC) or both national classification and IPC Applicant MEIJI SEIKA KAISHA, LTD. This opinion contains indications relating to the following items: Box No. I Basis of the opinion Box No. II Priority Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability Box No. IV Lack of unity of invention Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial Box No. V applicability; citations and explanations supporting such statement Box No. VI Certain documents cited Box No. VII Certain defects in the international application Box No. VIII Certain observations on the international application **FURTHER ACTION** 2. If a demand for international preliminary examination is made, this opinion will be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA") except that this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered. If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of 3 months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later. For further options, see Form PCT/ISA/220. For further details, see notes to Form PCT/ISA/220. 3. Name and mailing address of the ISA/JP Authorized officer Facsimile No. Telephone No.

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ł	Box	No. 1	Bas	s of this opinion		- 01/01/2003/003/07	
ŀ	1.	Witl					
		filed	, unless other	rise indicated under this item.	ished on the basis of the internation	onal application in the language in which it w	va
	ı	Ш	This opinion	has been established on the basis of a tr	ranslation from the original langua	ge into the following longue as	
		-		, which is the l	anguage of a translation furnished	for the purposes of international search (under	
			Rule 12.3 an	l 23.1(b)).		under the purposes of international search (under	er:
	2.	With inver	regard to an attion, this opin	y nucleotide and/or amino acid seq ion has been established on the basis of	uence disclosed in the internation	nal application and necessary to the claims	ed
		a.	type of mater	ai			
			a seque	nce listing			
			table(s)	related to the sequence listing			
		b.	format of mat	≠ial			
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		c.	time of filing/				
		ſ		d in the international application as filed	1		
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		Ī	•	subsequently to this Authority for the			
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3.	Ĺ	l f f	n addition, in urnished, the iled or does n	the case that more than one version o equired statements that the information it go beyond the application as filed, as	r copy of a sequence listing and/o in the subsequent or additional co appropriate, were furnished.	or table(s) relating thereto has been filed or opies is identical to that in the application as	
4.	A		onal comment				
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Box	No. I	V Lack of unity of invention
1.		In response to the invitation (Form PCT/ISA/206) to pay additional fees the applicant has:
		paid additional fees
		paid additional fees under protest
		not paid additional fees
2.	\boxtimes	This Authority found that the requirement of unity of invention is not complied with and chose not to invite the applicant to pay additional fees.
3.	This	Authority considers that the requirement of unity of invention in accordance with Rules 13.1, 13.2 and 13.3 is
		complied with
	\boxtimes	not complied with for the following reasons:
		The matter common to claims 1 to 9 and 10 to 12 resides in a β-fructofuranosidase mutant having a mutation in the amino acid sequence represented by SEQ ID NO:2.
		As the result of a search, however, it was clarified that such a β-fructofuranosidase mutant having a mutation in the amino acid sequence represented by SEQ ID NO:2 is not novel because of having been disclosed in documents "WO 97/34004 A1 (Meiji Seika Kaisha, Ltd.), 18 September, 1997 (18.09.97) & EP 889134 A1 & US 2002/0192771 A1".
		Therefore, this common matter falls within the category of the prior art and cannot be regarded as a special technical feature in the meaning within PCT Rule 13.2. Such being the case, there is no special technical matter common to all the claims and
		the aforesaid invention groups cannot be considered as a group of inventions so linked as to form a single general inventive concept.
		However, as all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
		•
4.	Con	sequently, this opinion has been established in respect of the following parts of the international application:
	\boxtimes	all parts
		the parts relating to claims Nos.

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Воз		Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement		
1.	Statement			
	Novelty (N)	Claims	1-16	YES
		Claims		NO NO
	Inventive step (IS)	Claims		YES
		Claims	1-16	NO
	Industrial applicability (L	A) Claims	1-16	YES
		Claims		NO

2. Citations and explanations:

Document 1: WO, 97-34004, A1 (Meiji Seika Kaisha, Ltd.), 18 September, 1997(18.09.97), & EP, 889134, A1, & US, 2002-0192771, A1

[Claims 1-16]

The subject matters of claims 1-16 do not appear to involve an inventive step in view of document 1 cited in the ISR.

Document 1 describes (1) a β -fructofuranosidase mutant derived from Aspergillus niger ACE-2-1(ATCC20611), (2) a DNA encoding the β -fructofuranosidase mutant, (3) an expression vector which comprises the DNA, (4) a host cell which comprises the expression vector, (5) a process for producing the β -fructofuranosidase mutant by cultivating the host cell, and (6) a process for producing a fructooligosaccharide comprising a step of bringing the β -fructofuranosidase mutant. Document 1 also describes (1) the mutant having the mutation in one amino acid residue (at positions 170, 300,313, or 386), (2) the mutant having the mutation in two amino acid residues (at positions 170+300, 300+313), and (3) the mutant having the mutation in three amino acid residues (at positions 170+300+313), as the β -fructofuranosidase mutant (see, claims 27-45, pages 46-66, and example D).

Furthermore, document 1 also describes that (1) the characteristics of β -fructofuranosidase with its amino acid sequence, and (2) 1-kestose can be produced selectively and efficiently using the β -fructofuranosidase mutant (see, page 6, lines 3-6 and page 7, lines 10-17). Document 1 also describes that the mutant having three amino acid residues to be mutated can produce 1-kestose more selectively than the mutant having one amino acid residue (for example, see, examples D1 and D6).

In addition, document 1 describes the amino acid sequences of the β -fructofuranosidase derived from Aspergillus niger, Penicillium roqueforti and Scopulariopsis brevicaulis, and the nucleotide sequence (see, SEQ ID No. 1, 2 and 11-14).

Document 1 describes that (1) 1-kestose and nystose make up part of the industrially produced fructooligosaccharide mixtures of today, and (2) their high-purity crystals exhibit new desirable characteristics both in physical properties and food processing purpose while maintaining the general physiological advantages of fructooligosaccharides (see, page 4, lines 12-18).

Considering that document 1 describes 1-kestose or nystose with high-purity has desirable characteristics, proposing to produce the 1-kestose or nystose with high-purity could have been easily conceived of. Furthermore, (1) creating a mutant by introducing a mutation randomly or site-specifically, (2) measuring desired characteristics, and (3) obtaining the mutant having the desired characteristics are well-known techniques to a person skilled in the art. So, when the 1-kestose or nystose with high-purity is produced, it is not considered to be especially difficult to (1) introduce a

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Box N	o. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
	nutation to the amino acid residues of the β-fructofuranosidase and (2) obtain a new mutant, in the process for producing the β-fructofuranosidase mutant described in document 1, and the effects of claims 1-16 could have been predicted from the description of document 1. Moreover, document 1 describes that the mutant having three amino acid residues to be nutated can produce 1-kestose more selectively than the mutant having one amino acid residue, and adging from this description, producing the β-fructofuranosidase mutant having mutations in amino acid residues at three positions could have been, as required, carried out.

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Box No. VIII

Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

The invention of the present application relates to a β -fructofuranosidase mutant. Claims 1, 4 and 10 show a plurality of the positions of the amino acid residue having a mutation, and claims 3, 6, 9 and 12 also show a plurality of substituted amino acids. However, the specification merely specifically discloses (1) the β -fructofuranosidase mutant (G62E, L122M, I128N, V165F, H221Y, Q395L, T550S, G40D, T381M, and W379C) having one substitution and (2) the β -fructofuranosidase mutant (V165F+G300V+H313K) having three substitutions, in the amino acid sequence represented by SEQ ID No. 2. So, the Claims are not supported by the specification as a whole.